

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
2 May 2002 (02.05.2002)

PCT

(10) International Publication Number  
**WO 02/34835 A1**

(51) International Patent Classification<sup>7</sup>: C08L 95/00, C08K 5/39, 9/08

(21) International Application Number: PCT/EP01/12341

(22) International Filing Date: 24 October 2001 (24.10.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:  
00 309 342.4 24 October 2000 (24.10.2000) EP

(71) Applicant (for all designated States except US): ATOFINA RESEARCH [BE/BE]; Société Anonyme, Zone Industrielle C, B-7181 Seneffe (Feluy) (BE).

(72) Inventor; and

(75) Inventor/Applicant (for US only): ANDRIOLI, Patrizio [BE/BE]; 14, rue des Mésanges, B-7140 Morlanwelz (BE).

(74) Common Representative: ATOFINA RESEARCH, S.A.; Patent Department, Zone Industrielle C, B-7181 Seneffe (Feluy) (BE).

(81) Designated States (national): AF, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

WO 02/34835 A1

(54) Title: BITUMEN VULCANISING COMPOSITION

(57) Abstract: Provided is a vulcanising agent for vulcanising bitumen, which vulcanising agent comprises a sulphur agent and a binder, wherein the vulcanising agent is in the form of pellets. Further provided is a process for producing a surfacing composition, which process comprises contacting bitumen with an elastomer and the vulcanising agent of the present invention.

## BITUMEN VULCANISING COMPOSITION

The present invention concerns a composition for vulcanising bitumen. The bitumen is then useful as a surfacing composition in, for example, road surfacing. The invention provides a vulcanising composition which when mixed with bitumen and an elastomer gives rise to a composition having improved stability and aging properties. The invention also relates to methods for forming the vulcanising composition and road surfacing composition, and to the surfacing composition itself.

It has been known for some time to add elastomeric polymers to bitumen to form a useful road surfacing composition. The elastomer provides the advantage of allowing the composition to deform under pressure (e.g. when a vehicle is driven across it) and then to return to its original state. This increases the lifetime of the road surface. It has also been known for some time to add sulphur or sulphur containing compounds to such compositions to further strengthen it by introducing cross-linking. This cross-linking is termed vulcanisation. The formation of such compositions is generally carried out in two steps: the elastomer is added to the bitumen and fully dispersed, and then the vulcanising agent is added and cross-linking is allowed to occur. A number of published patent applications describe such processes, including WO 98/47966, WO 92/11321, WO 90/02776, WO 93/18092, and WO 96/15193.

A specific process of this type is disclosed in published European patent, EP 424 420. This document discloses formation of a bitumen composition in two steps. An elastomeric polymer (SBS) is dispersed in bitumen at a temperature of around 140-180°C. When the SBS has fully dispersed in the bitumen, vulcanising agents are added, including sulphur, a sulphur donor and other additives. The mixture is agitated for from 100-150 minutes at a temperature of from 140-180°C to form the final composition.

In published French patent application FR 2 737 216 more recently developed an alternative one-step procedure for forming a surfacing composition from bitumen. This method makes use of a preliminary mixture ready for use, which preliminary mixture

comprises SBS in powder form and cross-linking agents in powder form. The preliminary mixture is itself thus a powder.

There are a number of disadvantages associated with all of the above methods. The two-step processes are more time consuming and require separate storage of a larger number of different components. Moreover, the elastomer and cross-linking agent are in powder form and must be added separately. Thus, their proportions need to be determined on forming the surfacing composition. This process is prone to error leading to wastage and increased cost. The one-step process overcomes some of these problems, but is itself associated with further problems. The preliminary mixture is in a powder form and needs to be mixed since it is composed of more than one ingredient. Powders are subject to safety problems, since they are a fire and explosion hazard. This is particularly problematic for sulphur and sulphur-containing compounds. Moreover, powders are difficult to store and are subject to clogging and caking which makes them difficult to process.

An aim of the present invention is to solve the problems associated with the above prior art. It is a further aim of the present invention to provide a vulcanising composition which is safer, easier to use and gives rise to a final surfacing composition which has improved stability, storage and aging characteristics and allows a better control on the amount of product used in the compositions.

Accordingly, the present invention provides a vulcanising agent for vulcanising bitumen, which vulcanising agent comprises a sulphur agent and a binder, wherein the vulcanising agent is in the form of pellets. Generally the sulphur agent is dispersed with the binder. By pellets, it is meant a product comprising particulate matter collected together to form larger agglomerates. These agglomerates are not especially limited in shape or size, provided that they are not in a hazardous powder form, and may include an extrudate, tablet or pill formed by extrusion or compression of the vulcanising agent.

The present invention provides a pelletised vulcanising agent which avoids the hazards associated with powders, and can be employed in a one-step process to form a road surfacing agent with an elastomer and bitumen. By one step process is meant a process in which the elastomer and vulcanising agent can be added together to the bitumen, if desired. A two-step process requires that the elastomer and bitumen are thoroughly dispersed before the vulcanising agent is added. Surprisingly, the present vulcanising agent also improves the aging and stability of the surfacing composition in addition to the above advantages.

The various aspects of the present invention will now be discussed in more detail.

The vulcanising agent comprises a binder and a sulphur agent, the proportions of which are not especially limited, provided that the binder is present in sufficient quantity to allow pellets to be formed. In a preferred embodiment the vulcanising agent comprises 20 wt.% or more of the binder. More preferably the vulcanising agent comprises from 20-90 wt.% of the binder and from 10-80 wt.% of the sulphur agent. The vulcanising agent may also comprise from 0-30 wt.% of further additives. Preferably the sulphur agent comprises from 10-100 wt.% of elemental sulphur and from 0-90 wt.% of a sulphur-containing compound. The sulphur agent may thus comprise a sulphur donor, such as zinc dibutyl dithiocarbamate (ZDBC) or zinc mercaptobenzothiazol (ZMBT) or a sulphinamide. Preferred sulphur containing-compounds and sulphur donors may be found in the Rubber Handbook published by the Swedish Institution of Rubber Technology.

The binder is not especially limited, provided that it is capable of pelletising the sulphur agent. Preferably, however, the binder has a melting point (and/or softening point) below the melting point of sulphur, for ease of processing. In preferred embodiments, the binder comprises a wax, a hydrocarbon resin, a copolymer of ethylene and an acrylic ester. The more preferred binders include polyethylene (PE), glycol monostearate (GMS), ethyl vinyl alcohol (EVA) and copolymers of ethylene and acrylate esters.

A particularly preferred vulcanising agent is elemental sulphur, and a particularly preferred binder is EVA.

The vulcanising agent preferably comprises further additives. These additives are not especially limited and may be added to improve mixing and/or facilitate processing. They may also be selected to improve surface quality, especially road quality. Such further additives may comprise tackifiers, elastomers, bitumens, zinc oxide, and/or stearic acid.

The present invention also provides a process for production of the vulcanising agent, which process comprises:

- (a) extruding a sulphur agent with a binder; or
- (b) compressing a sulphur agent with a binder,

at a temperature below the melting temperature of the sulphur agent, so as to form pellets of the vulcanisation agent.

The process is a standard compression or extrusion process and is well known in the art. Standard processing techniques and devices may be employed. The process preferably employs a temperature of 110°C or less, since sulphur melts at around 119°C. However, if the sulphur agent as a whole has a higher melting temperature than this, then higher temperatures may be employed.

The present invention further provides a process for producing a surfacing composition, which process comprises contacting bitumen with an elastomer and a vulcanising agent as defined above. In a preferred embodiment, the process is carried out at a temperature of 100°C or more. More preferably the process is carried out at a temperature of 120-200°C. As already mentioned above, this is a one-step process where the elastomer and vulcanising agent are added substantially simultaneously to the bitumen.

Generally from 1-15 parts by weight of elastomer are employed with from 85-99 parts by weight of bitumen. Preferably from 0.01-10 wt.%, more preferably from 0.1-5 wt.% of vulcanising agent is employed, based upon the total weight of bitumen and elastomer.

The elastomer is not especially limited provided that it has the qualities required for surfacing compositions. Such elastomers are well known in the art and are generally rubbery polymers. In a preferred embodiment, the elastomer comprises styrene butadiene styrene (SBS), hydrogenated SBS, styrene isoprene styrene (SIS), styrene ethylene butadiene styrene (SEBS) and/or polyisobutadiene (PIB).

The surfacing composition of the present invention has improved aging and stability properties and is preferably used for producing and/or repairing a road, pavement or track, or other surface for vehicles.

Typically such surfacing compositions are made up prior to transporting them to the site at which they are to be used. Once made up they are stored for up to three days at temperatures of around 180°C, to keep the bitumen from solidifying. Thus, stability at high temperatures is important for such compositions, and it is important that the compositions undergo as little phase separation as possible. The present surfacing compositions are especially stable, as shown in the ring and ball test (described below) as compared with prior art compositions.

The present invention will now be described with reference to the following specific embodiments, which are by way of Example only and are not intended to limit the invention.

### Examples

In the following examples, the bitumens employed were a Venezuelan bitumen (Bitumen A) having a penetration of 87 at 25°C, and a Middle Eastern bitumen (Bitumen B) having a penetration of 74 at 25°C. Additionally five Chinese bitumens have been tested. They have penetration values at 25 °C of 75, 58, 78, 83 respectively, for

SH1, SH2, SH3 and SH4. The penetration test is a standard well known test in the industry, in which the depth of penetration of a needle into the bitumen is measured. It was measured here following the methods of standard test ASTM D-5-73 revised 95. All these bitumen were used for laboratory tests.

The elastomer employed in the laboratory tests was Finaprene® 503 (SBS - a 31/69 wt.% styrene/butadiene polymer). The elastomers employed in the industrial road compositions were Finaprene® 503 (SBS - a linear 31/69 wt.% styrene/butadiene polymer), Finaprene® 401 (a radial low molecular weight 20/80 wt% styrene/butadiene polymer) and Finaprene® 411X (a radial high molecular weight 30/70 wt% styrene/butadiene polymer)

The vulcanising agent comprised either:

- (a) sulphur powder,
- (b) 70/30 wt.% of sulphur/EVA pellets prepared by extrusion; or
- (c) 70/30 wt.% of (80/20 wt.% sulphur/ZDBC)/EVA pellets prepared by compression.
- (d) 30/62/3/3/2 wt% of EVA/sulphur/ZDBC/ZMBT/PIB pellets prepared by extrusion

#### Example 1.

This is an example of the present invention.

Bitumen A was heated to 180°C and 3 wt.% of Finaprene® 503 was added in pellet form, simultaneously with 0.1 wt.% vulcanisation agent © in pellet form, based on the total weight of bitumen and elastomer. After 120 minutes the mixture was homogeneous. The stability to storage was good (there was no phase separation) and confirmed the efficacy of the vulcanisation.

**Example 2.**

This is an example of the present invention.

Bitumen B was heated to 180°C and 5 wt.% of Finaprene® 503 was added in pellet form, simultaneously with 0.1 wt.% vulcanisation agent (b) in pellet form, based on the total weight of bitumen and elastomer. After 120 minutes the mixture was homogeneous. The stability to storage was good (there was no phase separation) and confirmed the efficacy of the vulcanisation.

**Example 3.**

This is a comparative example employing a method of FR 2 737 216.

Bitumen B was heated to 180°C and 5 wt.% of Finaprene® 503 was added in powder form, simultaneously with 0.1 wt.% vulcanisation agent (a) in powder form, based on the total weight of bitumen and elastomer. After 120 minutes the mixture was homogeneous. The stability to storage was less good than for Examples 1 and 2.

**Stability and aging test using the ring and ball method.**

The products of Example 2 and Comparative Example 3 were tested for stability. An internal standard method developed from the ring and ball method (ASTM-D36) was employed. The surfacing composition is coated on a ring after it has been made, then the ring is submerged in hot oil. The temperature at which the composition falls away from the ring is recorded. The composition is stored at high temperature for 3 days and the method is repeated. If the stability is good, the composition will retain its tackiness and the temperature will ideally remain as close to the originally measured temperature as possible. The larger the drop in temperature, the lower the stability of the composition and the worse its aging characteristics.

The results for the compositions formed in Examples 2 and 3 are shown below in Table 1.

**Table 1**

Mixing time / mins	Drop in temp / °C	
	Example 2	Example 3
120	7	10
180	5	10

Table 1 shows that for mixing times of 2 and 3 hours, there is a lesser drop in temperature for the composition of the present invention compared with the closest prior art composition. This highlights the improvement in aging and stability characteristics of the present compositions.

**Examples 4 to 7.**

In these examples, several surfacing compositions were prepared from the Chinese Bitumens SH1, SH2, SH3 and SH4, with various elastomers in different amounts and optionally with various amounts of heavy furfural extract. The compositions according to the present invention were prepared with 0.1 wt% of vulcanising agent d) in pellet form and the comparative examples were prepared without vulcanising agent. The compositions and results are displayed in Tables 2 to 5. In these examples, the ductility was measured following the method of standard test IP 32/55, the kinematic viscosity of asphalt was measured following the method of standard test ASTM D 2170 and the elastic recovery was measured following the method of standard test DIN V 52021-1.

**Table 2.**  
**Bitumen SH1.**

Composition	wt%	wt%	wt%
Bitumen SH1	96.5	96.4	95.5
Finaprene® 503	-	-	-
Finaprene® 401	3.5	3.5	4.5
Vulcanising agent	-	0.1	-
Heavy furfural extract	-	-	-
Properties			
Ring and ball temperature- °C	68	70	80
Penetration 5°C - 1/10 mm	10	11	13
15 °C - 1/10 mm	18	20	18
25 °C - 1/10 mm	51	55	54
Viscosity 135 °C - Pa.s	1.22	1.47	1.54
150 °C - Pa.s	0.72	0.84	0.89
Ductility 5 °C - cm	13	14	20
13 °C - cm	60	80	60
25 °C - cm	62	87	58
Elastic recovery - %	92	95	97
Storage stability (48 hrs at 163 °C)	no	yes	no

**Table 3.****Bitumen SH2.**

Composition	wt%	wt%	wt%	wt%	wt%
Bitumen SH2	97.5	97.4	96.5	96.4	95.5
Finaprene® 503	-	-	-	-	-
Finaprene® 401	2.5	2.5	3.5	3.5	4.5
Vulcanising agent	-	0.1	-	0.1	-
Heavy furfural extract	-	-	-	-	-
Properties					
Ring and ball temperature -°C	53	55	67	69	74
Penetration 5 °C - 1/10 mm	9	10	9	10	9
15 °C - 1/10 mm	16	10	15	17	16
25 °C - 1/10 mm	47	44	43	51	44
Viscosity 135 °C - Pa.s	1.16	1.16	1.31	1.52	1.76
150 °C - Pa.s	0.63	0.70	0.79	0.85	0.98
Ductility 5 °C - cm	6	1	8	8	14
13 °C - cm	24	41	52	58	60
25 °C - cm	97	>130	54	82	53
Elastic recovery - %	35	58	93	93	98
Storage stability (48 hrs at 163 °C)	no	yes	no	yes	No

Table 4.  
Bitumen SH3.

Composition	Wt%											
Bitumen SH3	96.5	96.4	95.5	97.5	97.4	97.5	97.5	96.5	96.4	96.5	96.5	95.5
Finaprene® 503	3.5	3.5	4.5	-	-	-	-	-	-	-	-	-
Finaprene® 401	-	-	-	2.5	2.5	2.5	2.5	3.5	3.5	3.5	3.5	4.5
Vulcanising agent	-	0.1	-	-	0.1	-	-	0.1	-	-	-	-
Heavy furfural extract	-	-	-	-	-	5%	10%	-	-	5%	10%	-
Properties												
Ring and ball temperature -°C	52	54	93	50	52	48	45	70	72	65	58	76
Penetration 5°C - 1/10 mm	13	12	11	11	10	-	-	10	12	-	-	10
15 °C - 1/10 mm	22	23	22	21	20	-	-	21	21	-	-	21
25 °C - 1/10 mm	57	60	53	66	60	88	124	58	57	74	110	53
Viscosity 135 °C - Pa.s	1.03	1.12	1.28	0.89	0.99	0.86	0.67	1.18	1.30	0.95	0.79	1.51
150 °C - Pa.s	0.63	0.66	0.79	0.55	0.59	-	0.67	0.78	-	-	-	0.82
Ductility 5 °C - cm	20	32	39	11	11	22	57	22	19	46	77	26
13 °C - cm	71	73	102	37	54	-	-	49	74	-	-	66
25 °C - cm	>140	>140	103	110	>130	-	-	52	65	-	-	52
Elastic recovery - %	68	68	98	28	60	65	58	98	95	95	95	100
Storage stability (48 hrs at 163 °C)	no	yes	no	yes	Yes	yes	yes	no	yes	yes	yes	no

**Table 5.****Bitumen SH4.**

Composition	wt%	wt%	wt%	wt%	wt%	wt%
Bitumen SH4	97	96.9	96.5	96.4	96	95.9
Finaprene® 503	-	-	-	-	-	-
Finaprene® 401	3	3	3.5	3.5	4	4
Vulcanising agent	-	0.1	-	0.1	-	0.1
Properties						
Ring and ball Temperature -°C	57	59	73	74	80	82
Penetration 5 °C - 1/10 mm	10	12	14	11	11	11
15 °C - 1/10 mm	23	20	22	23	19	19
25 °C - 1/10 mm	68	57	43	6	58	55
Viscosity 135 °C - Pa.s	1.00	1.16	1.09	1.10	1.2	1.47
150 °C - Pa.s	0.61	0.65	0.72	0.77	0.74	0.86
Ductility 5 °C - cm	15	21	16	41	25	27
13 °C - cm	46	51	70	96	72	84
25 °C - cm	54	109	55	7	56	70
Elastic recovery - %	78	73	95	98	98	98
Storage stability (48 hrs at 163 °C)	yes	yes	no	yes	no	Yes

The results displayed in Tables 2 to 5 clearly show that all the bitumen compositions prepared according to the present inventions had excellent storage stability. In addition, as desired, the ring and ball temperature was higher for the compositions of the present invention. The viscosity was also higher for the compositions using the vulcanising agent of the invention resulting in a better cohesion and so was the ductility. It must be observed in Table 4 that a higher ductility can be obtained by the addition of 5 to 10 wt% of heavy furfural extract, but at the expense of the ring and ball temperature that is lower and at the expense of the penetration value that is higher.

**Industrial Examples.**

Two road compositions have been prepared with the Chinese bitumen SH2, the elastomer Finaprene® 503 and the vulcanising composition d) in pellet form and in powder form

respectively. The small drop in temperature observed for the laboratory tests of Examples 1 and 2 was confirmed for the industrial examples. This can be seen in Table 6 showing the drop in ring and ball temperature as a function of ageing for the same vulcanising agent in pellet form and in powder form.

**Table 6.**

**Difference in Ring and Ball Temperature after ageing.**

Vulcanising agent	Pellets	Powder
$\Delta T$ °C		
1 day at 180 °C	2.3	5.6
2 days at 180 °C	3.3	6.0
3 days at 180 °C	3.3	5.7

## CLAIMS:

1. A vulcanising agent for vulcanising bitumen, which vulcanising agent comprises a sulphur agent and a binder, wherein the vulcanising agent is in the form of pellets.
2. A vulcanising agent according to claim 1, comprising:
  - (i) 20-90 wt.% of the binder; and
  - (ii) 10-80 wt.% of the sulphur agent; and
  - (iii) 0-30 wt.% of additives.
3. A vulcanising agent according to claim 1 or claim 2, wherein the sulphur agent comprises from 10-100 wt.% of elemental sulphur and from 0-90 wt.% of a sulphur-containing compound.
4. A vulcanising agent according to any preceding claim, wherein the sulphur agent comprises a sulphur donor.
5. A vulcanising agent according to claim 4, wherein the sulphur donor comprises zinc dibutyl dithiocarbamate (ZDBC).
6. A vulcanising agent according to any preceding claim, wherein the binder comprises a wax, a hydrocarbon resin, a copolymer of ethylene and an acrylic ester, polyethylene (PE), glycol monostearate (GMS) and/or ethyl vinyl alcohol (EVA).
7. A vulcanising agent according to any preceding claim, wherein the additives comprise a tackifier, an elastomer, bitumen, zinc oxide and/or stearic acid.

15.

8. A vulcanising agent according to any preceding claim, wherein the pellets are in the form of an extrudate, tablet or pill formed by extrusion or compression of the vulcanising agent.

9. A process for production of a vulcanising agent, which process comprises:

- (a) extruding a sulphur agent with a binder; or
- (b) compressing a sulphur agent with a binder,

at a temperature below the melting temperature of the sulphur agent, so as to form pellets of the vulcanisation agent.

10. A process according to claim 9, wherein the temperature is a temperature of 110°C or less.

11. A process according to claim 9 or claim 10, wherein the vulcanising agent comprises an agent as defined in any of claims 2-8.

12. A vulcanising agent obtainable according to a process as defined in any of claims 9-11.

13. A process for producing a surfacing composition, which process comprises contacting bitumen with an elastomer and a vulcanising agent as defined in any of claims 1-8 and 12.

14. A process according to claim 13, wherein the contacting is carried out at a temperature of 100°C or more.

15. A process according to claim 14, wherein the contacting is carried out at a temperature of 120-200°C.

16. A process according to any of claims 13-15, wherein from 1-10 parts by weight of elastomer are employed with from 90-99 parts by weight of bitumen.
17. A process according to any of claims 13-16 wherein from 0.01-10 wt.% of vulcanising agent is employed, based upon the total weight of bitumen and elastomer.
18. A process according to any of claims 13-17, wherein the elastomer comprises styrene butadiene styrene (SBS), hydrogenated SBS, styrene isoprene styrene (SIS), styrene ethylene butadiene styrene (SEBS) and/or polyisobutadiene (PIB).
19. A surfacing composition obtainable according to a process as defined in any of claims 13-18.
20. Use of the vulcanising agent in the form of pellets of the present invention to reduce health hazards.
20. Use of a surfacing composition as defined in claim 19 for producing and/or repairing a road, pavement or track, or other surface for vehicles.
21. Use of a surfacing composition as defined in claim 19 in roofing compositions and membranes, waterproof bitumen compositions membranes or mastics, joint sealants, or sound deadening or sound proofing compositions or membranes.

## INTERNATIONAL SEARCH REPORT

Intel Application No  
PCT/EP 01/12341

A. CLASSIFICATION OF SUBJECT MATTER  
IPC 7 C08L95/00 C08K5/39 C08K9/08

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
IPC 7 C08L C08K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

WPI Data, EPO-Internal

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5 795 929 A (GRUBBA WILLIAM E) 18 August 1998 (1998-08-18) column 1, paragraph 1 claims 1,2,14 ---	1-3,12, 13,17,18
A	US 6 025 418 A (DEFOOR FRANS MAURICE JOSEPH ET AL) 15 February 2000 (2000-02-15) examples ---	1,5,13, 18
A	EP 0 384 254 A (BASF AG) 29 August 1990 (1990-08-29) claims 1,8; example 1 ---	1,4,13, 18
A	GB 2 270 318 A (BASF CORP) 9 March 1994 (1994-03-09) examples; table V ---	1,4,13, 18,19 19
X		-/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

## \* Special categories of cited documents :

- \*A\* document defining the general state of the art which is not considered to be of particular relevance
- \*E\* earlier document but published on or after the international filing date
- \*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*O\* document referring to an oral disclosure, use, exhibition or other means
- \*P\* document published prior to the international filing date but later than the priority date claimed

\*T\* later document published after the International filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

\*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

\*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

\*&\* document member of the same patent family

Date of the actual completion of the international search

Date of mailing of the international search report

19 February 2002

01/03/2002

Name and mailing address of the ISA

Authorized officer

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2290 HV Rijswijk  
Tel (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax: (+31-70) 340-3016

Engel, H

## INTERNATIONAL SEARCH REPORT

International Application No  
PCT/EP 01/12341

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 360 656 A (ELF FRANCE) 28 March 1990 (1990-03-28) cited in the application the whole document	1-5, 13-19

## INTERNATIONAL SEARCH REPORT

Information on patent family members

Int'l Application No  
PCT/EP 01/12341

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
US 5795929	A	18-08-1998	NONE		
US 6025418	A	15-02-2000	DE DE DK EP FI NO RO RU	68922613 D1 68922613 T2 287890 A 0424420 A1 98823 B 301239 B1 108352 B1 2011666 C1	14-06-1995 05-10-1995 08-02-1991 02-05-1991 15-05-1997 29-09-1997 28-04-1994 30-04-1994
EP 0384254	A	29-08-1990	EP	0384254 A2	29-08-1990
GB 2270318	A	09-03-1994	CA	2104908 A1	27-02-1994
EP 0360656	A	28-03-1990	FR AT AU AU BR DE DE DK EP ES WO JP JP NO US	2636340 A1 99723 T 628390 B2 4217689 A 8907085 A 68912044 D1 68912044 T2 113690 A 0360656 A1 2062069 T3 9002776 A1 2731442 B2 3501035 T 178580 B 5508112 A	16-03-1990 15-01-1994 17-09-1992 02-04-1990 02-10-1990 17-02-1994 11-08-1994 28-06-1990 28-03-1990 16-12-1994 22-03-1990 25-03-1998 07-03-1991 15-01-1996 16-04-1996